

ATROPINE SULFATE- atropine sulfate injection, solution

Vedco, Inc

Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, click [here](#).

Atropine Sulfate

NDC 50989-075-12

ATROPINE SULFATE

Injection 1/120 Grain

STERILE MULTIPLE DOSE VIAL

NET CONTENTS: 100 mL

VEDCO

KEEP OUT OF REACH OF CHILDREN

FOR ANIMAL USE ONLY

CAUTION: Federal law restricts this drug to use by or on the order of a licensed veterinarian.

For Intravenous, Intramuscular, or Subcutaneous Use

COMPOSITION: Each mL contains:

Atropine Sulfate 0.54 mg

Sodium Chloride 9 mg

Benzyl Alcohol (preservative)..... 1.5%

Water for Injection q.s.

pH adjusted with sulfuric acid when necessary.

DOSAGE AND ADMINISTRATION:

Dogs and Cats: Inject 1 mL for each 20 lbs. of body weight as a pre-anesthetic adjuvant, or to reduce salivation, bronchial secretions, or internal peristalsis associated with colic or diarrhea.

As an antidote for parasympathomimetic drugs, 1 mL for each 7.5 lbs. of body weight. It is suggested that 1/4 of the dosage be injected intravenous and the remainder intramuscular or subcutaneous.

WARNING: Poisonous alkaloid. Keep out of reach of children

Antidotes: warmth, emetics, cholinergics.

Store at room temperature between 15° and 30°C (59°- 86°F)

Rev. 3/08

Distributed By

VEDCO, INC

St. Joseph, MO 64507

TAKE TIME OBSERVE LABEL DIRECTIONS

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Lot No./Exp. Date:

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Product Information

Product Type	PRESCRIPTION ANIMAL DRUG	Item Code (Source)	NDC:50989-075
Route of Administration	INTRAVENOUS, INTRAMUSCULAR, SUBCUTANEOUS		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ATROPINE SULFATE (UNII: 03J5ZE7KA5) (ATROPINE - UNII:7C0697DR9I)	ATROPINE SULFATE	0.54 mg in 1 mL

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:50989-075-12	100 mL in 1 VIAL, MULTI-DOSE		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		03/01/2008	

Labeler - Vedco, Inc (021634266)

Revised: 6/2018

Vedco, Inc